

NATALIZUMAB PRIOR AUTHORIZATION FORM (form effective 1/5/2026)

Prior authorization guidelines for **Natalizumab** and **Quantity Limits/Daily Dose Limits** are available on Geisinger Health Plan's website at <https://healthplan.geisinger.org/pharmacy/pharmacy.aspx?strip=true&style=OneGeisinger>.

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		total pages: _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:		NPI:	State license #:	
LTC facility contact/phone:		Street address:		
Beneficiary name:		City/state/zip:		
Beneficiary ID#:	DOB:	Phone:	Fax:	

CLINICAL INFORMATION

Drug requested:	Strength:	Quantity: _____ vials	Refills:
Directions: <input type="checkbox"/> 300 mg SQ every 4 weeks <input type="checkbox"/> other: _____			
Diagnosis (<u>submit documentation</u>):		Dx code (<u>required</u>):	
Is the beneficiary currently being treated with the requested medication?	<input type="checkbox"/> Yes – date of last dose: _____ <i>Submit documentation.</i> <input type="checkbox"/> No		
Is natalizumab prescribed by or in consultation with a neurologist or gastroenterologist?	<input type="checkbox"/> Yes <i>Submit documentation of consultation if applicable.</i> <input type="checkbox"/> No		
Is the beneficiary receiving chronic immunosuppressive or immune modulating therapies?	<input type="checkbox"/> Yes <i>Submit complete medication list.</i> <input type="checkbox"/> No		

Complete all sections that apply to the beneficiary and this request.
Check all that apply and submit documentation for each item.

INITIAL requests

<p>1. For treatment of MULTIPLE SCLEROSIS (MS):</p> <p><input type="checkbox"/> Has a relapsing form of MS</p>
<p>2. For treatment of CROHN'S DISEASE (CD):</p> <p><input type="checkbox"/> Has moderate-to-severe CD</p> <p><input type="checkbox"/> Has CD that is associated with high-risk or poor prognostic features</p> <p><input type="checkbox"/> Has achieved remission with the requested medication AND:</p> <p style="padding-left: 20px;"><input type="checkbox"/> Will be using the requested medication as maintenance therapy to maintain remission</p> <p><input type="checkbox"/> Tried and failed a TNF-inhibitor (e.g., Cimzia, Humira, Remicade) or has a contraindication or an intolerance to TNF-inhibitors</p> <p><input type="checkbox"/> Tried and failed an IL-12/23 or IL-23 inhibitor (e.g., Skyrizi, Stelara, Tremfya) or has a contraindication or an intolerance to IL-12/23 and IL-23 inhibitors</p> <p><input type="checkbox"/> Tried and failed or has a contraindication or intolerance to vedolizumab (Entyvio)</p>

3. For a NON-PREFERRED natalizumab product:

- Has a history of trial and failure of or a contraindication or an intolerance to the preferred natalizumab product(s) approved or medically accepted for the beneficiary's diagnosis

RENEWAL requests

1. For treatment of MULTIPLE SCLEROSIS (MS):

- Experienced improvement or stabilization of the MS disease course since starting natalizumab

2. For treatment of CROHN'S DISEASE:

- Experienced therapeutic benefit within 3 months of starting natalizumab
 Was able to discontinue concomitant steroid use within 6 months of starting natalizumab (if applicable)
 Has been using natalizumab for at least 1 year AND:
 Has not required additional steroid use for disease control for more than 3 months in the past 12 months

3. For a NON-PREFERRED natalizumab product:

- Has a history of trial and failure of or a contraindication or an intolerance to the preferred natalizumab product(s) approved or medically accepted for the beneficiary's diagnosis

Please submit to PromptPA <https://ghp.promptpa.com>

OR fax to Geisinger Health Plan at 570-271-5610 the completed form with required clinical documentation.

Prescriber Signature:

Date:

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